## **AMENDMENTS TO THE CLAIMS**

This listing of claims will replace all prior versions, and listings, of claims in the application:

- (Original) A tissue comprising linkers bonded to the tissue and a bridge molecule bonded between two or more of the linkers, wherein the linkers and the bridges are chemically different.
- 2. (Original) The tissue of claim 1 wherein the tissue comprises extracellular matrix selected from the group consisting of collagenous fibrils, GAG and elastin.
- 3. (Original) The tissue of claim 1 wherein the two linkers and the bridge bonded between the two linkers span a distance of between about 10 Angstroms and about 100 Angstroms.
- 4. (Original) The tissue of claim 1 wherein the two linkers and the bridge bonded between the two linkers span a distance of between about 15 Angstroms and about 50 Angstroms.
  - 5. (Original) The tissue of claim 1 wherein the bridge is a single molecule.
- 6. (Original) The tissue of claim 1 wherein the bridge is reactive with modified tissue.

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7. (Original) The tissue of claim 1 wherein the bridge comprises functional groups

selected from the group consisting of methylthio, thio, amine, alcohol, carboxyl and

combinations thereof.

8. (Original) The tissue of claim 1 wherein the bridge comprises a hydrocarbon

backbone.

9. (Original) The tissue of claim 1 wherein the linkers comprise monomers, dimers

and oligomers.

10. (Currently Amended) The tissue of claim 1 wherein the linkers are active

substantially nonreactive with respect to the unmodified tissue.

11. (Original) The tissue of claim 1 wherein the linkers comprise functional groups

selected from the group consisting of aldehydes, epoxies, imide groups, photooxidative

groups, enzymatically oxidative groups and combinations thereof.

12. (Original) The tissue of claim 1 wherein the linkers comprise crosslinking

agents.

13. (Original) The tissue of claim 1 wherein the linker is selected from the group

consisting of glutaraldehyde, triglycidyl amine and epoxy.

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14. (Original) The tissue of claim 1 wherein a bioprosthetic device comprises the tissue.

15. (Original) The tissue of claim 14 wherein the bioprosthetic device is a heart

valve prosthesis.

16. (Original) A method of crosslinking tissue comprising treating the tissue with a

linker composition comprising linkers and a bridge composition comprising bridges wherein

the linkers bond to the tissue and the bridges bond between two of the linkers, wherein the

bridges and the linkers are chemically different.

17. (Original) The method of claim 16 wherein the tissue comprises proteins.

------18. (Original)- The method of claim 16-wherein the tissue is treated with the linker

composition and the bridge composition simultaneously.

19. (Original) The method of claim 16 wherein the tissue is treated with the linker

composition prior to addition of the bridge composition.

20. (Original) The method of claim 16 wherein the linker composition and the

bridge composition are combined prior to treating the tissue.

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Docket Number: 01610.0048-US-01 Office Action Response 21. (Original) The method of claim 16 wherein the linker composition comprises

crosslinking agents.

22. (Original) The method of claim 16 wherein the concentration of the linkers in

the linker composition is between about 0.0001 molar and about molar.

23. (Original) The method of claim 16 wherein the concentration of the bridges in

the bridge composition is between about  $1 \times 10^{-7}$  molar and about 1 molar.

24. (Original) The method of claim 16 wherein the tissue is treated with the linker

composition and the bridge composition for between about 10 minutes and about one

month.

25. (Original) The method of claim 16-wherein the tissue is treated with the linker

composition and the bridge composition for between about 10 minutes and about 2 weeks.

26. (Original) The method of claim 16 wherein the bridges comprise multiple

functional groups.

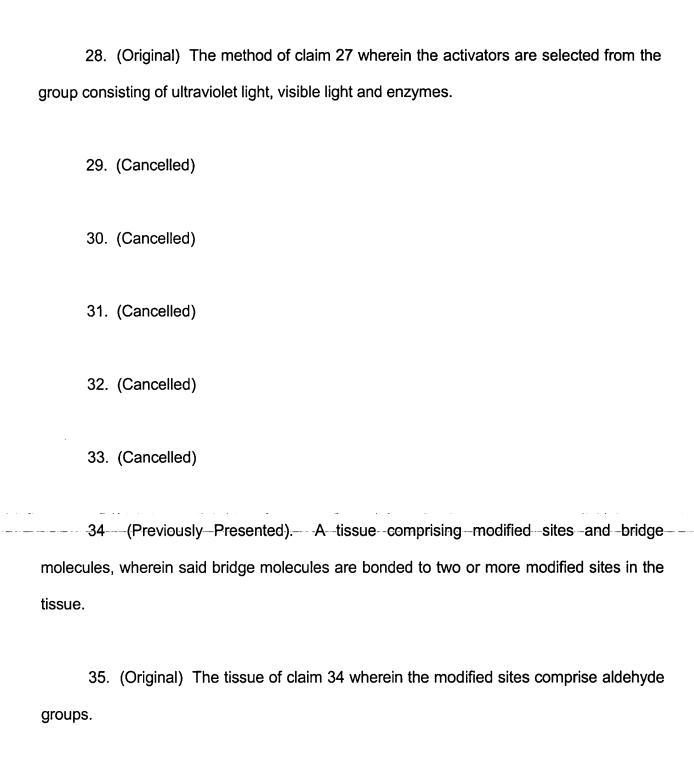
27. (Original) The method of claim 16 wherein the treatment of the tissue further

comprises exposing the tissue to activators.

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wherein the bridge molecules bond to two or more modified sites in the tissue.

36. (Previously Presented) A method of crosslinking tissue having modified sites

comprising treating said tissue with a bridge composition comprising bridge molecules

- 37. (Original) The method of claim 36 wherein the modified sites comprises aldehyde groups.
- 38. (Previously Presented) The tissue of claim 34 wherein said bridge molecules comprise functional groups selected from the group consisting of methylthio, amine, alcohol, carboxyl and combinations thereof.
- 39. (Previously Presented) The tissue of claim 34 wherein said bridge molecules are substantially non-reactive to tissues having no modified sites.
- 40. (Previously Presented) The tissue of claim 34 wherein said bridge molecules comprise functional groups located at opposite ends of said bridge molecules.
- 41. (Previously Presented) The method of claim 36 wherein said bridge molecules comprise functional groups selected from the group consisting of methylthio, amine, alcohol, carboxyl and combinations thereof.
- 42. (Previously Presented) The method of claim 36 wherein said bridge molecules comprise a functional groups located at opposite ends of said bridge molecules.

43. (Previously Presented) The tissue of claim 1 wherein said bridge molecules are substantially non-reactive to tissues having no modified sites.